

's Docket No.: 07039-170002

TES PATENT AND TRADEMARK OFFICE

COPY OF PAPERS

Applicant: Jorg J. Goronzy et al.

Art Unit : 1648 **ORIGINALLY FILED**

Serial No.: 09/723,000

Examiner: Stacy Brown

Filed

: November 27, 2000

Title

: METHODS AND MATERIALS FOR EVALUATING RHEUMATOID

ARTHRITIS

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Commissioner for Patents Washington, D.C. 20231

RESPONSE TO NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE AND/OR AMINO ACID SEQUENCES

In response to the communication dated July 2, 2002 (copy enclosed), Applicants respectively submit that no sequence listing is required since (1) all sequence listing requirements were complied with as explained in Applicants' Communication mailed June 4, 2002 and (2) claims 58 and 59 have been cancelled as set forth in the Preliminary Amendment filed herewith.

Please apply any charges or credits to Deposit Account No. 06-1050.

Respectfully submitted,

atrick Finn III, Ph.D. Reg. No. 44,109

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CERTIFICATE OF MAILING BY FIRST CLASS MAIL

I hereby certify under 37 CFR §1.8(a) that this correspondence is being deposited with the United States Postal Service as first class mail with sufficient postage on the date indicated below and is addressed to the Commissioner for Patents, Washington, D.C. 20231.

Date of Deposit

Signature

Dasilku

Typed or Printed Name of Person Signing Certificate

Application N .: 09/723,000

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. § 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application	n does not
comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821	 1.825 for the
following reason(s):	

	1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to these regulations, published at 1114 OG 29, May 15, 1990 and at 55 FR 18230, May 1, 1990.	
	 This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c). 	
	3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).	
	4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."	
	5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).	
	6. The paper copy of the "Sequence Listing" is not the same as the computer readable form of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).	
	7. Other: <u>Claims 58-59 require SEQ 10 NO; they contain</u> references to amino acid positions of a polypeptide	
Applicant Must Provide:		
M	An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".	
Ø	An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.	
X	A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).	
For	questions regarding compliance to these requirements, please contact: Rules Interpretation, call (703) 308-4216 CRF Submission Help, call (703) 308-4212 tentIn Software Program Support (SIRA) Technical Assistance	

PLEASE RETURN A COPY OF THIS NOTICE WITH YOUR RESPONSE